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Substitute for form 1449/PTO  <b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b>  (Use as many sheets as necessary)			<b>Complete if Known</b>		
			Application Number	10/650,449-Conf. #7321	
			Filing Date	August 27, 2003	
			First Named Inventor	Michael L. Robinson	
			Art Unit	1649	
			Examiner Name	O. Chernyeshev	
Sheet	1	of	3	Attorney Docket Number	28335/39524A

U.S. PATENT DOCUMENTS					
Examiner Initials*	Cite No. <sup>1</sup>	Document Number Number-Kind Code <sup>2</sup> (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear

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NON PATENT LITERATURE DOCUMENTS				
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	C1	Accession No. AC003964, "Homo Sapiens Chromosome 16 BAC CIT987SK-A-911E12," (2002).		
	C2	Accession No. AC027281, "Homo Sapiens Chromosome 16 Clone RP11-424M24," (2003).		
	C3	Accession No. AC069308, "Mus Musculus Strain C57BL6/J Chromosome 8 RPC123-21B7," (2002).		
	C4	Accession No. AC109135, "Homo Sapiens Chromosome 1 Clone RP11-240C17," (2003).		
	C5	Accession No. AC130459, "Homo Sapiens Chromosome 1 Clone CTA-427H10," (2006).		
	C6	AL-SHROOF et al., "Ciliary Dyskinesia Associated with Hydrocephalus and Mental Retardation in a Jordanian Family," <i>Mayo Clin. Proc.</i> , 76(12): 1219-1224 (2001).		
	C7	BANNISTER, "Some Scanning Electron Microscopic Observations of the Ependymal Surface of the Ventricles of Hydrocephalic Hy3 Mice and a Human Infant," <i>Acta Neurochir</i> , 46:159-168 (1979).		
	C8	BERRY, "The Inheritance and Pathogenesis of Hydro-Cephalus-3 in the Mouse," <i>J. Path. Bact.</i> , 81:157-161 (1961).		
	C9	BLAKE et al., "The Mouse Genome Database (MGD): The Model Organism Database for the Laboratory Mouse," <i>Nucl. Acids Res.</i> , 30: 113-115 (2002).		
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	C11	CALLEN et al., "Re-Evaluation of GM2346 From a Del(16)(q22) to T(4;16)(q35;q22.1)," <i>Clin. Genet.</i> , 38:466-468 (1990).		
	C12	CASTRO-GAGO et al., "Autosomal Recessive Hydrocephalus with Aqueductal Stenosis," <i>Childs Nerv. Syst.</i> 12: 188-191 (1996).		
	C13	CHOW et al., "Autosomal Recessive Hydrocephalus with Third Ventricle Obstruction," <i>Am. J. Med Genet</i> , 35:310-313 (1990).		
	C14	CHUDLEY et al., "Bilateral Sensorineural Deafness and Hydrocephalus Due to Foramen of Monro Obstruction in Sibs: A Newly Described Autosomal Recessive Disorder," <i>Am. J. Med Genet.</i> , 68: 350-360 (1997).		

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C15	CLARK, "Hydrocephalus, A Hereditary Character in the House Mouse," <i>Proc. Natl. Acad. USA</i> , 18:654-656 (1932).	
C16	DAHME et al., "Disruption of the Mouse <i>L1</i> Gene Leads to Malformations of the Nervous System," <i>Nature Genetics</i> , 17:346-349 (1997).	
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C18	DAVIES et al., "A Detailed Investigation of Two Cases Exhibiting Characteristics of the 6p Deletion Syndrome," <i>Hum. Genet.</i> , 98: 454-459 (1996).	
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C23	FRANSEN et al., "L1-Associated Diseases: Clinical Geneticists Divide, Molecular Geneticists Unite," <i>Hum. Mol. Genet.</i> 6:1625-1632 (1997).	
C24	FRANSEN et al., "L1 Knockout Mice Show Dilated Ventricles, Vermis Hypoplasia and Impaired Exploration Patterns," <i>Hum. Mol. Genet.</i> , 7: 999-1009 (1998).	
C25	FRYNS et al., "Interstitial 16q Deletion with Typical Dysmorphic Syndrome," <i>Ann Genet.</i> , 24:124-125 (1981).	
C26	GALBREATH et al., "Overexpression of TGF- $\beta$ 1 in the Central Nervous System of Transgenic Mice Results in Hydrocephalus," <i>J. Neuropathol. Exp. Neurol.</i> , 54: 339-49 (1995).	
C27	GAME et al., "Fetal Growth Retardation, Hydrocephalus, Hypoplastic Multilobed Lungs, and Other Anomalies in 4 Sibs," <i>Am. J. Med Genet.</i> , 33: 276-279 (1989).	
C28	GRÜNEBERG, "Congenital Hydrocephalus in the Mouse, a Case of Spurious Pleiotropism," <i>J. Genet.</i> , 45: 1-28 (1943).	
C29	HOLLANDER, "Hydrocephalic-Polydactyl, a Recessive Pleiotropic Mutant in the Mouse, and its Location in Chromosome 6," <i>Iowa State J. Res.</i> , 51: 13-23 (1976).	
C30	HOMANICS et al., "Targeted Modification of the Apolipoprotein B Gene Results in Hypobetalipoproteinemia and Developmental Abnormalities in Mice," <i>Proc. Natl. Acad. Sci. USA</i> , 90: 2389-2393 (1993).	
C31	HUANG et al., "apo B Gene Knockout in Mice Results in Embryonic Lethality in Homozygotes and Neural Tube Defects, Male Infertility, and Reduced HDL Cholesterol Ester and apo A-I Transport Rates in Heterozygotes," <i>J. Clin. Invest.</i> 96: 2152-2161 (1995).	
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C36	MCNEISH et al., "Phenotypic Characterization of the Transgenic Mouse Insertional Mutation, <i>Legless</i> ," <i>J. Exp. Zool.</i> , 253: 151-162 (1990).	
C37	MELTON, "Gene Targeting in the Mouse," <i>Bioassays</i> , 16: 633-638 (1994).	
C38	MILLONIG et al., "The Mouse <i>Dreher</i> Gene <i>Lmx1a</i> Controls Formation of the Roof Plate in the	

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		Vertebrate CNS," <i>Nature</i> , 403: 764-769 (2000).	
	C39	NARITOMI et al., "16q21 is Critical for 16q Deletion Syndrome," <i>Clin. Genet.</i> , 33:372-375 (1988).	
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